



## ABCG2 gene

ATP binding cassette subfamily G member 2 (Junior blood group)

### Normal Function

The *ABCG2* gene belongs to a group of genes called the ATP-binding cassette family; genes in this family provide instructions for making proteins that transport molecules across cell membranes. In the intestines, the *ABCG2* protein helps release (secrete) a substance called urate into the urine. Urate is a byproduct of certain normal biochemical reactions in the body. In the bloodstream it acts as an antioxidant, protecting cells from the damaging effects of unstable molecules called free radicals. Urate levels are regulated by the kidneys and, to a lesser extent, by the intestines.

The *ABCG2* protein also transports certain drugs out of cells. For example, this protein clears some chemotherapy drugs from organs and tissues. Transport of these drugs allows them to have their intended effects and be eliminated from the body.

### Health Conditions Related to Genetic Changes

#### Gout

Genetic changes in the *ABCG2* gene have been found to be associated with a condition called gout, which is a form of arthritis that causes painful joint inflammation.

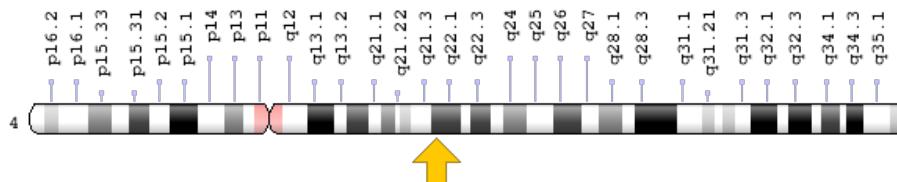
*ABCG2* gene changes associated with gout decrease the protein's ability to release urate. One variant replaces the protein building block (amino acid) glutamine with the amino acid lysine at position 141 in the protein (written as Gln141Lys or Q141K). This change reduces the protein's ability to secrete urate by half. Another variant creates a premature stop signal in the instructions for making the *ABCG2* protein (written as Gln126Ter or Q126X), which results in no functional *ABCG2* protein. Variants in the *ABCG2* gene reduce the removal of urate in the blood, which causes the blood level of urate to rise. The excess urate can accumulate in the body's joints in the form of crystals, triggering an inflammatory response from the immune system and leading to gout.

While changes in the *ABCG2* gene can alter urate levels in the body, they are not enough to cause gout by themselves. A combination of dietary, genetic, and other environmental factors play a part in determining the risk of developing this complex disorder.

## Chromosomal Location

Cytogenetic Location: 4q22.1, which is the long (q) arm of chromosome 4 at position 22.1

Molecular Location: base pairs 88,090,264 to 88,231,626 on chromosome 4 (Homo sapiens Updated Annotation Release 109.20200522, GRCh38.p13) (NCBI)



Credit: Genome Decoration Page/NCBI

## Other Names for This Gene

- ABC15
- ABCP
- ATP-binding cassette transporter G2
- ATP-binding cassette, sub-family G (WHITE), member 2 (Junior blood group)
- BCRP
- BCRP1
- BMDP
- breast cancer resistance protein
- CD338
- CDw338
- EST157481
- mitoxantrone resistance-associated protein
- MRX
- multi drug resistance efflux transport ATP-binding cassette sub-family G (WHITE) member 2
- MXR
- MXR-1
- MXR1
- placenta-specific ATP-binding cassette transporter

- placenta specific MDR protein
- UAQTL1

## **Additional Information & Resources**

### Educational Resources

- Biochemistry (fifth edition, 2002): Purines Are Degraded to Urate in Human Beings  
<https://www.ncbi.nlm.nih.gov/books/NBK22372/#A3526>
- Informed Health Online: Gout: Overview  
<https://www.ncbi.nlm.nih.gov/books/NBK284934/>

### Scientific Articles on PubMed

- PubMed  
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28ABCG2%5BTI%5D%29+OR+%28ATP+binding+cassette+subfamily+G+member+2%5BTI%5D%29%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1080+days%22%5Bdp%5D>

### Catalog of Genes and Diseases from OMIM

- ATP-BINDING CASSETTE, SUBFAMILY G, MEMBER 2  
<http://omim.org/entry/603756>

### Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology  
[http://atlasgeneticsoncology.org/Genes/GC\\_ABCG2.html](http://atlasgeneticsoncology.org/Genes/GC_ABCG2.html)
- ClinVar  
<https://www.ncbi.nlm.nih.gov/clinvar?term=ABCG2%5Bgene%5D>
- HGNC Gene Symbol Report  
[https://www.genenames.org/data/gene-symbol-report/#!/hgnc\\_id/HGNC:74](https://www.genenames.org/data/gene-symbol-report/#!/hgnc_id/HGNC:74)
- Monarch Initiative  
<https://monarchinitiative.org/gene/NCBIGene:9429>
- NCBI Gene  
<https://www.ncbi.nlm.nih.gov/gene/9429>
- UniProt  
<https://www.uniprot.org/uniprot/Q9UNQ0>

## Sources for This Summary

- OMIM: ATP-BINDING CASSETTE, SUBFAMILY G, MEMBER 2  
<http://omim.org/entry/603756>
- Köttgen A, Albrecht E, Teumer A, Vitart V, Krumsiek J, Hundertmark C, Pistis G, Ruggiero D, O'Seaghdha CM, Haller T, Yang Q, Tanaka T, Johnson AD, Katalik Z, Smith AV, Shi J, Struchalin M, Middelberg RP, Brown MJ, Gaffo AL, Pirastu N, Li G, Hayward C, Zemunik T, Huffman J, Yengo L, Zhao JH, Demirkan A, Feitosa MF, Liu X, Malerba G, Lopez LM, van der Harst P, Li X, Kleber ME, Hicks AA, Nolte IM, Johansson A, Murgia F, Wild SH, Bakker SJ, Peden JF, Dehghan A, Steri M, Tenesa A, Lagou V, Salo P, Mangino M, Rose LM, Lehtimäki T, Woodward OM, Okada Y, Tin A, Müller C, Oldmeadow C, Putku M, Czamara D, Kraft P, Frogheri L, Thun GA, Grotevendt A, Gislason GK, Harris TB, Launer LJ, McArdle P, Shuldiner AR, Boerwinkle E, Coresh J, Schmidt H, Schallert M, Martin NG, Montgomery GW, Kubo M, Nakamura Y, Tanaka T, Munroe PB, Samani NJ, Jacobs DR Jr, Liu K, D'Adamo P, Uliivi S, Rotter JI, Psaty BM, Vollenweider P, Waeber G, Campbell S, Devuyst O, Navarro P, Kolcic I, Hastie N, Balkau B, Froguel P, Esko T, Salumets A, Khaw KT, Langenberg C, Wareham NJ, Isaacs A, Kraja A, Zhang Q, Wild PS, Scott RJ, Holliday EG, Org E, Viigimaa M, Bandinelli S, Metter JE, Lupo A, Trabetti E, Sorice R, Döring A, Lattka E, Strauch K, Theis F, Waldenberger M, Wichmann HE, Davies G, Gow AJ, Bruinenberg M; LifeLines Cohort Study, Stolk RP, Kooner JS, Zhang W, Winkelmann BR, Boehm BO, Lucae S, Penninx BW, Smit JH, Curhan G, Mudgal P, Plenge RM, Portas L, Persico I, Kirin M, Wilson JF, Mateo Leach I, van Gilst WH, Goel A, Ongen H, Hofman A, Rivadeneira F, Uitterlinden AG, Imboden M, von Eckardstein A, Cucca F, Nagaraja R, Piras MG, Nauck M, Schurmann C, Budde K, Ernst F, Farrington SM, Theodoratou E, Prokopenko I, Stumvoll M, Jula A, Perola M, Salomaa V, Shin SY, Spector TD, Sala C, Ridker PM, Kähönen M, Viikari J, Hengstenberg C, Nelson CP; CARDIoGRAM Consortium; DIAGRAM Consortium; ICBP Consortium; MAGIC Consortium, Meschia JF, Nalls MA, Sharma P, Singleton AB, Kamatani N, Zeller T, Burnier M, Attia J, Laan M, Klopp N, Hillege HL, Kloiber S, Choi H, Pirastu M, Tore S, Probst-Hensch NM, Völzke H, Gudnason V, Parsa A, Schmidt R, Whitfield JB, Fornage M, Gasparini P, Siscovick DS, Polasek O, Campbell H, Rudan I, Bouatia-Naji N, Metspalu A, Loos RJ, van Duijn CM, Borecki IB, Ferrucci L, Gambaro G, Deary IJ, Wolffenbuttel BH, Chambers JC, März W, Pramstaller PP, Snieder H, Gyllensten U, Wright AF, Navis G, Watkins H, Witteman JC, Sanna S, Schipf S, Dunlop MG, Tönjes A, Ripatti S, Soranzo N, Toniolo D, Chasman DI, Raitakari O, Kao WH, Ciullo M, Fox CS, Caulfield M, Bochud M, Gieger C. Genome-wide association analyses identify 18 new loci associated with serum urate concentrations. *Nat Genet.* 2013 Feb;45(2):145-54. doi: 10.1038/ng.2500. Epub 2012 Dec 23.  
*Citation on PubMed:* <https://www.ncbi.nlm.nih.gov/pubmed/23263486>  
*Free article on PubMed Central:* <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3663712/>
- Matsuo H, Takada T, Ichida K, Nakamura T, Nakayama A, Takada Y, Okada C, Sakurai Y, Hosoya T, Kanai Y, Suzuki H, Shinomiya N. Identification of ABCG2 dysfunction as a major factor contributing to gout. *Nucleosides Nucleotides Nucleic Acids.* 2011 Dec;30(12):1098-104. doi: 10.1080/15257770.2011.627902.  
*Citation on PubMed:* <https://www.ncbi.nlm.nih.gov/pubmed/22132963>
- Matsuo H, Takada T, Nakayama A, Shimizu T, Sakiyama M, Shimizu S, Chiba T, Nakashima H, Nakamura T, Takada Y, Sakurai Y, Hosoya T, Shinomiya N, Ichida K. ABCG2 dysfunction increases the risk of renal overload hyperuricemia. *Nucleosides Nucleotides Nucleic Acids.* 2014; 33(4-6):266-74. doi: 10.1080/15257770.2013.866679.  
*Citation on PubMed:* <https://www.ncbi.nlm.nih.gov/pubmed/24940678>
- Merriman T. Genomic Influences on Hyperuricemia and Gout. *Rheum Dis Clin North Am.* 2017 Aug; 43(3):389-399. doi: 10.1016/j.rdc.2017.04.004. Review.  
*Citation on PubMed:* <https://www.ncbi.nlm.nih.gov/pubmed/28711141>

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